



Guidelines for the Use of Antiretroviral Agents in Pediatric HIV Infection

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Table 15d. Antiretroviral Therapy-Associated Adverse Effects and Management Recommendations—Hematologic Effects (Last updated April 14, 2020; last reviewed April 14, 2020) (page 1 of 2)

| Adverse Effects | Associated ARVs | Onset/Clinical Manifestations | Estimated Frequency | Risk Factors | Prevention/Monitoring | Management |
|---------------------------|-----------------|--|--|---|--|--|
| Anemia^a | ZDV | <p>Onset:</p> <ul style="list-style-type: none"> Variable; weeks to months after starting therapy <p>Presentation</p> <p><i>More Common:</i></p> <ul style="list-style-type: none"> Asymptomatic Mild fatigue Pallor Tachypnea <p><i>Rare:</i></p> <ul style="list-style-type: none"> Congestive heart failure | <p>Newborns Exposed to HIV:</p> <ul style="list-style-type: none"> Severe anemia is uncommon, but may be seen coincident with physiologic Hgb nadir. <p>Children with HIV Who Are Taking ARV Drugs:</p> <ul style="list-style-type: none"> Anemia is two to three times more common with ZDV-containing regimens than with all other regimens. | <p>Newborns Exposed to HIV:</p> <ul style="list-style-type: none"> Premature birth is the most common risk factor <i>In utero</i> exposure to ZDV-containing regimens Advanced maternal HIV Neonatal blood loss Combination ARV prophylaxis or presumptive HIV therapy, particularly ZDV plus 3TC <p>Children with HIV Who Are Taking ARV Drugs:</p> <ul style="list-style-type: none"> Underlying hemoglobinopathy (e.g., sickle cell disease, G6PD deficiency) Myelosuppressive drugs (e.g., TMP-SMX, rifabutin) Iron deficiency Advanced or poorly controlled HIV disease OIs of the bone marrow Malnutrition | <p>Newborns Exposed to HIV:</p> <ul style="list-style-type: none"> Obtain CBC at birth. Consider repeating CBC at 4 weeks for neonates who are at higher risk (e.g., those born prematurely or who are known to have low birth Hgb) and for neonates who receive ZDV beyond 4 weeks. <p>Children with HIV Who Are Taking ARV Drugs:</p> <ul style="list-style-type: none"> Avoid using ZDV in children with severe anemia when alternative agents are available. Obtain CBC as part of routine care (see Clinical and Laboratory Monitoring of Pediatric HIV Infection). | <p>Newborns Exposed to HIV:</p> <ul style="list-style-type: none"> Anemia rarely requires intervention unless it is symptomatic or Hgb <7.0 g/dL. ZDV administration can be limited to 4 weeks in low-risk neonates (see Antiretroviral Management of Newborns with Perinatal HIV Exposure or HIV Infection). <p>Children with HIV Who Are Taking ARV Drugs:</p> <ul style="list-style-type: none"> Discontinue non-ARV, marrow-toxic drugs, if feasible. Treat coexisting iron deficiency, OIs, and malignancies. For persistent, severe anemia that is thought to be associated with ARV drugs (typically macrocytic anemia), switch to a regimen that does not contain ZDV. |
| Macrocytosis | ZDV | <p>Onset:</p> <ul style="list-style-type: none"> Within days or weeks of starting therapy <p>Presentation:</p> <ul style="list-style-type: none"> Asymptomatic, but MCV is often >100 fL Sometimes associated with anemia | >90% to 95% for all ages | None | No monitoring required—macrocytosis can be detected if CBC is obtained as part of routine care (see Clinical and Laboratory Monitoring of Pediatric HIV Infection). | No management required. |

Table 15d. Antiretroviral Therapy-Associated Adverse Effects and Management Recommendations—Hematologic Effects (Last updated April 14, 2020; last reviewed April 14, 2020) (page 2 of 2)

| Adverse Effects | Associated ARVs | Onset/Clinical Manifestations | Estimated Frequency | Risk Factors | Prevention/Monitoring | Management |
|--------------------------|-----------------|--|---|---|--|--|
| Neutropenia ^a | ZDV | Onset: <ul style="list-style-type: none"> • Variable Presentation: <ul style="list-style-type: none"> • Asymptomatic | Newborns Exposed to HIV: <ul style="list-style-type: none"> • Rare Children with HIV Who Are Taking ARV Drugs: <ul style="list-style-type: none"> • 2% to 4% of children on ARV drugs • Highest rates occur in children on ZDV-containing regimens | Newborns Exposed to HIV: <ul style="list-style-type: none"> • <i>In utero</i> exposure to ARV drugs • Combination ARV prophylaxis, particularly ZDV plus 3TC Children with HIV Who Are Taking ARV Drugs: <ul style="list-style-type: none"> • Advanced or poorly controlled HIV infection • Myelosuppressive drugs (e.g., TMP-SMX, ganciclovir, hydroxyurea, rifabutin) | Children with HIV Who Are Taking ARV Drugs: <ul style="list-style-type: none"> • Obtain CBC as part of routine care. | Newborns Exposed to HIV: <ul style="list-style-type: none"> • No established threshold for intervention; some experts would consider using an alternative NRTI for prophylaxis if ANC reaches <500 cells/mm³. ZDV administration can be limited to 4 weeks in low-risk neonates (see Antiretroviral Management of Newborns with Perinatal HIV Exposure or HIV Infection). Children with HIV Who Are Taking ARV Drugs: <ul style="list-style-type: none"> • Discontinue non-ARV, marrow-toxic drugs, if feasible. • Treat coexisting OIs and malignancies. • In cases of persistent, severe neutropenia that is thought to be associated with ARV drugs, switch to a regimen that does not contain ZDV. |

^a HIV infection itself, OIs, and medications that are used to prevent OIs (e.g., TMP-SMX) may all contribute to anemia and neutropenia.

Key: 3TC = lamivudine; ANC = absolute neutrophil count; ARV = antiretroviral; CBC = complete blood count; dL = deciliter; fL = femtoliter; G6PD = glucose-6-phosphate dehydrogenase; Hgb = hemoglobin; MCV = mean cell volume; NRTI = nucleoside reverse transcriptase inhibitor; OI = opportunistic infection; TMP-SMX = trimethoprim-sulfamethoxazole; ZDV = zidovudine

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